



ORIGINAL ARTICLE

Early amplitude-integrated electroencephalography for monitoring neonates at high risk for brain injury[☆]



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KEYWORDS

Newborn;
Brain injury;
Amplitude-integrated
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Early outcome

Abstract

Objective: This study aimed to correlate amplitude-integrated electroencephalography findings with early outcomes, measured by mortality and neuroimaging findings, in a prospective cohort of infants at high risk for brain injury in this center in Brazil.

Methods: This blinded prospective cohort study evaluated 23 preterm infants below 31 weeks of gestational age and 17 infants diagnosed with hypoxic-ischemic encephalopathy secondary to perinatal asphyxia, with gestational age greater than 36 weeks, monitored with amplitude-integrated electroencephalography in a public tertiary center from February 2014 to January 2015. Background activity (classified as continuous, discontinuous high-voltage, discontinuous low-voltage, burst-suppression, continuous low-voltage, or flat trace), presence of sleep-wake cycling, and presence of seizures were evaluated. Cranial ultrasonography in preterm infants and cranial magnetic resonance imaging in infants with hypoxic-ischemic encephalopathy were performed.

Results: In the preterm group, pathological trace or discontinuous low-voltage pattern ($p=0.03$) and absence of sleep-wake cycling ($p=0.019$) were associated with mortality and brain injury assessed by cranial ultrasonography. In patients with hypoxic-ischemic encephalopathy, seizure patterns on amplitude-integrated electroencephalography traces were associated with mortality or brain lesion in cranial magnetic resonance imaging ($p=0.005$).

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PALAVRAS-CHAVE

Recém-nascido;
Lesão cerebral;
EEG de amplitude
integrada;
Resultado precoce

Conclusion: This study supports previous results and demonstrates the utility of amplitude-integrated electroencephalography for monitoring brain function and predicting early outcome in the studied groups of infants at high risk for brain injury.

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Eletoencefalograma de amplitude integrada precoce no monitoramento de neonatos com risco elevado de lesão cerebral

Resumo

Objetivo: Este estudo visou correlacionar os achados do eletroencefalograma de amplitude integrada (aEEG) com resultados precoces, medidos por mortalidade e achados de neuroimagem, em uma coorte prospectiva de neonatos com risco elevado de lesão cerebral em nosso centro no Brasil.

Métodos: O estudo prospectivo de coorte cego avaliou 23 neonatos prematuros abaixo de 31 semanas de idade gestacional (IG) e 17 neonatos diagnosticados com Encefalopatia Hipóxico-Isquêmica (EHI) secundária à asfíxia perinatal, com IG superior a 36 semanas, monitorados com aEEG em um centro terciário público de fevereiro de 2014 a janeiro de 2015. Foram avaliadas a atividade de base (classificada como padrão contínuo, descontínuo de alta voltagem, descontínuo de baixa voltagem, supressão de explosão, contínuo de baixa voltagem ou traço plano), a presença de ciclo do sono-vigília e a presença de convulsões. Foram feitas a ultrassonografia craniana em prematuros e a ressonância magnética (RM) craniana em neonatos com EHI.

Resultados: No grupo de prematuros, o traço patológico ou padrão descontínuo de baixa voltagem ($p = 0,03$) e a ausência de ciclo do sono-vigília ($p = 0,019$) foram associados a mortalidade e lesão cerebral avaliada por ultrassonografia craniana. Em pacientes com EHI, os padrões de convulsão nos traçados do aEEG foram associados a mortalidade ou lesão cerebral na RM craniana ($p = 0,005$).

Conclusão: Este estudo corrobora os resultados anteriores e demonstra a utilidade do aEEG no monitoramento da função cerebral e na predição de alterações precoces nos grupos de neonatos estudados com risco elevado de lesão cerebral.

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Introduction

The incidence of neurodevelopmental impairment in extremely preterm infants and those with hypoxic-ischemic encephalopathy (HIE) secondary to perinatal asphyxia remains high in spite of advances in perinatal care. Studies have estimated a global incidence of 345,000 premature infants and 233,000 infants with HIE per year with moderate/severe neurological impairment.^{1,2} Both populations are considered at high risk for brain injury.

Different imaging methods can evaluate brain injury and assess neurological prognosis.³⁻⁵ Acute electroencephalographic abnormalities result from neuronal disorganization and correlate with cognitive impairment.^{6,7} Amplitude-integrated electroencephalography (aEEG) provides a clinically accessible method for continuous observation of cerebral background activity in ill infants at the bedside. Thus, the utility of early aEEG for the assessment of the severity of cerebral injury and adverse outcomes in premature infants has been investigated as a tool for assessing initial neonatal risk.⁸⁻¹⁰

The aEEG pattern is well correlated with conventional EEG, and results in term newborns with perinatal asphyxia

showed good predictive value of short- and long-term neurological prognosis.¹¹⁻¹³ Other studies have shown that severe electroencephalographic abnormalities in preterm neonates evaluated during the first 72 h of life are related to neurodevelopmental impairment.^{14,15}

Given the severity of brain injury, as well its high morbidity and mortality rate, the identification of prognostic factors with appropriate timing to provide early future interventions is relevant. Therefore, this study aimed to correlate aEEG findings with early outcomes, measured by mortality and neuroimaging findings, in a prospective cohort of infants at high risk for brain injury in this center in Brazil.

Methods

This study was performed in a public tertiary center in Brazil. All infants born between February 2014 and January 2015, with gestational age (GA) above 36 weeks with HIE secondary to perinatal asphyxia or with GA below 31 weeks, were prospectively included in the study after parental consent was obtained. All subjects included were inborn. The study was approved by the institutional ethics committee. Infants with genetic syndromes or congenital malformations

incompatible with life were excluded, as these conditions could affect the results.

The criteria for perinatal hypoxic-ischemic events included the presence of at least two of the following: (a) Apgar score below 5 at 5 min; (b) need for ventilation up to 10 min of life (intubation or ventilation under continuous positive airway pressure); (c) gas analysis of cord blood in the first hour of life with $\text{pH} < 7.10$ or $\text{BE} > -12$. The modified clinical Sarnat score was used to grade HIE.¹⁶ The aEEG was recorded for 72 h as a two-channel EEG from biparietal surface disk electrodes using an EEG device (Neuron-Spectrum-4 and 5 systems with Neurospectrum software module for aEEG and trending – Neurosoft, Russia). Newborns with HIE who were treated with the institutional hypothermia protocol were evaluated for an additional 24 h in order to monitor the rewarming phase. In brief, the obtained signal was filtered, rectified, smoothed, and amplitude-integrated before it was printed out or digitally available on the monitor at a slow speed (6 cm/h), directly at the bedside.

The recordings were analyzed by two independent readers and registered as shown below¹⁷:

- 1) Background activity:
 - Continuous voltage pattern (CVP): continuous activity with minimum amplitude above $5 \mu\text{V}$ and maximum amplitude above $10 \mu\text{V}$.
 - Discontinuous voltage pattern (DC): discontinuous activity with minimum amplitude below $5 \mu\text{V}$ and maximum amplitude above $10 \mu\text{V}$. For preterm evaluation this study also used the sub-classification proposed by Olischar et al.¹⁸ and this pattern was further subdivided into: (a) discontinuous high voltage pattern (DHVP) – minimum range between 3 and $5 \mu\text{V}$; (b) discontinuous low voltage pattern (DLVP) – minimum range below $3 \mu\text{V}$.
 - Burst suppression: discontinuous activity with minimum amplitude without variability at $0-1/2 \mu\text{V}$, and bursts with amplitude $>25 \mu\text{V}$.
 - Inactive, flat trace (FT): mainly inactive background (electrocerebral inactivity) with amplitude always below $5 \mu\text{V}$.
- 2) Sleep-wake cycling (SWC): characterized by sinusoidal smooth cyclic variations, mainly of the minimum amplitude. It was further categorized as developed SWC, immature SWC, and no SWC.
- 3) Seizures: characterized as an abrupt rise in minimum and maximum amplitude, categorized as single seizures (no more than one seizure per each period of 30 min of analysis), repetitive seizures (more than one electrographic seizure per each 30 min period of analysis but no more than one electrographic seizure over a 10 min period), or status epilepticus (ongoing seizure activity >30 min, present as “sawtooth pattern” or as continuous increase of the lower and upper margins, or more than one electrographic seizure over a 10 min period).¹⁹
- 4) Newborns with HIE were further assessed by time to normal trace (TTNT): calculated by the number of hours of life to regain normal aEEG activity (continuous voltage pattern).

To characterize background activity, 4-h periods of adequate monitoring were evaluated daily on days 1, 2, and 3 of life. Both aEEG readers were blinded to the clinical history. Cranial ultrasound (cUS) was performed at least once a week in the neonatal period and cranial magnetic resonance imaging (MRI) was performed in all infants with HIE between days four and ten of life. Clinical and imaging data were collected including with pre-established interest variables, such as demographic (sex, maternal age, race), clinical (birth weight, gestational age, Apgar scores, diagnosis of early-onset sepsis, presence of clinical seizure, need for ventilatory support, surfactants, and vasoactive drugs), and imaging data. The MRI examinations were assessed by two independent neuroradiologists without knowledge of the clinical data or aEEG findings. Since the interpretation agreement among different MRI raters may affect the reliability of results, the interobserver agreement was also evaluated in this study. MRI was performed with T1 spin echo sequences; T1 MTC; T2 FSE; diffusion; and T2 gradient-echo. Four radiographic findings were evaluated: lesions in the posterior limb of the internal capsule (PLIC) – labeled as present or absent; lesions in the basal ganglia and thalamus (BGT), white matter, and cortical gray matter, labeled as normal, mild, moderate, or severe.

In the premature group, good early outcome was defined as survival to 28 days without severe periventricular/intraventricular hemorrhage (PIVH), defined as grade III or IV, or peri-ventricular leukomalacia (PVL), while poor early outcome was considered as neonatal death or presence of severe PIVH or PVL. In newborns from the HIE group, good early outcome was defined as survival without moderate/severe lesion assessed by cranial MRI, while poor early outcome was considered as death or moderate/severe lesion assessed by cranial MRI. Presence of PIVH was evaluated according to Papile classification. The HIE cohort pattern of brain injury was classified according to abnormalities in brain regions known to be susceptible in HIE, based on criteria published by Rutherford et al.²⁰

Statistical analysis

A descriptive analysis was performed using percentage and number of valid cases for categorical variables as well as mean, standard deviation, and number of valid cases for continuous variables in order to identify the main characteristics of patients with high risk for brain injury monitored with aEEG. Fisher’s exact test was used for analysis of categorical variables and the *t*-test for comparisons of continuous variables using Stata software (Stata Corp. 2013, Stata Statistical Software, version 13, USA), considering a 5% significance level for all tests. Positive predictive value (PPV), negative predictive value (NPV), and risk ratio (RR) were evaluated. For inter-rater agreement analysis, the kappa coefficient (κ) was applied, using the SPSS v. 19.0 software.

Results

Patients

During the study period, a total of 40 patients were included; 23 extremely premature neonates and 17 with HIE. In

Table 1 Baseline characteristics of monitored groups.

Extremely preterm group	Total (n=23)	Good early outcome (n=10)	Poor early outcome (n=13)	p value
Gestational age (wk) (mean ± SD)	28 ± 1.7	28.3 ± 1.3	27.6 ± 1.6	NS
Weight (g) (mean ± SD)	917 ± 243	965 ± 261	880 ± 185	NS
Male (n)	12	6	6	NS
Apgar 1 (mean ± SD)	5.2 ± 2.5	5.9 ± 2.8	4.7 ± 2.1	NS
Apgar 5 (mean ± SD)	7.9 ± 2.6	8.9 ± 1.2	7.1 ± 3.2	NS
HIE group	Total (n=17)	Good early outcome (n=11)	Poor early outcome (n=6)	p value
Gestational age (wk) (mean ± SD)	39 ± 1.5	39.3 ± 1	38.6 ± 2.2	NS
Weight (g) (mean ± SD)	3100 ± 422	3139 ± 217	3029 ± 680	NS
Male (n)	14	8	6	NS
Apgar 1 (mean ± SD)	2.2 ± 1	2.2 ± 0.9	2.3 ± 1.3	NS
Apgar 5 (mean ± SD)	4.7 ± 1.3	4.8 ± 1	4.7 ± 1.8	NS
Sarnat score (n)				
Mild encephalopathy	0	0	0	NS
Moderate encephalopathy	10	6	4	NS
Severe encephalopathy	7	5	2	NS
Therapeutic hypothermia (n)	17	11	6	NS

HIE, hypoxic-ischemic encephalopathy; wk, weeks; SD, standard deviation.

the group of extremely preterm infants, gestational age ranged from 26 to 30.4 weeks, with a mean of 28 weeks; birth weight ranged from 610g to 1310g, with a mean of 938g. Eight infants in this group died during the study period. In the HIE group, gestational age ranged from 36 to 40.6 weeks, with a mean of 39 weeks; birth weight ranged from 2280g to 3940g, with a mean of 3100g, and one infant died. The baseline characteristics are shown in Table 1.

aEEG findings

All patients started monitoring within the first day of life (mean 6.2h of life, range 2–16h of life) and were monitored for 72–96h after birth. Pathological aEEG trace was defined as flat tracing, burst-suppression, and continuous low-voltage pattern.

Extremely preterm newborns

In the preterm group, mean time to initiate aEEG monitoring was 5.2 (±1.2) h of life, and pathological trace or DLVP were associated with higher rates of poor early outcome ($p=0.03$, PPV 90.9%, NPV 75%, RR = 3.63). Absence of SWC was associated with poor early outcome ($p=0.019$, PPV 75%, NPV 87.5%, RR = 1.53). The results are shown in Table 2. The background activity of patients with poor early outcome changed from normal to pathological aEEG pattern in five patients (38.4%), while in one (7.6%) it changed from pathological to normal aEEG pattern. None of the patients with good early outcome developed pathological aEEG pattern during the period of monitoring. Evolution of aEEG background activity in patients with poor early outcome and good early outcome is shown in Fig. 1.

Newborns with HIE

Patients with HIE were assessed for background activity in the first hours of life, TTNT, presence of SWC, and seizures. Seizure patterns on aEEG traces ($p=0.005$, PPV 83.3%, NPV 90.9%, RR = 5.45) and longer TTNT ($p=0.015$) were associated with poor early outcome. All the patients with HIE underwent therapeutic hypothermia. The results are shown in Table 2.

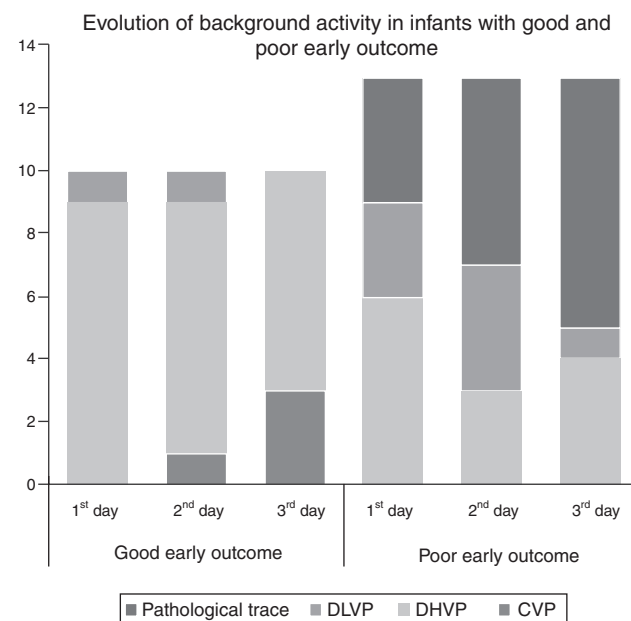


Figure 1 Evolution of the background activity over the first 3 days of life in preterm infants with good and poor early outcomes. DLVP, discontinuous low-voltage pattern; DHVP, discontinuous high-voltage pattern; CVP, continuous voltage pattern.

Table 2 aEEG findings related to outcome in extremely preterm and HIE groups.

Extremely preterm group	Total (n = 23)	Good early outcome (n = 10)	Poor early outcome (n = 13)	p value
<i>Background activity</i>				
Pathological trace	9	0	9	0.002
DLVP	9	1	8	0.029
Pathological trace or DLVP	11	1	10	0.003
DHVP or continuous	12	9	3	NS
<i>SWC</i>				
Developed	0	0	0	0.019
Immature	7	6	1	
Absent	16	4	12	
HIE group	Total (n = 17)	Good early outcome (n = 11)	Poor early outcome (n = 6)	p value
<i>aEEG background activity, 3–6 h of life</i>				
Continuous	11	9	2	NS
Not continuous	6	2	4	NS
<i>TTNT (mean)</i>	12.17	5.36	24.7	0.015
<i>Presence of SWC</i>	11	9	2	NS
<i>Seizures (n)</i>				
Single seizures	0	0	0	0.005
Repetitive seizures	3	1	2	NS
Status epilepticus	3	0	3	0.029
Repetitive or status epilepticus	6	1	5	0.005
Clinical seizures	4	1	3	NS

aEEG, amplitude-integrated electroencephalography; HIE, hypoxic-ischemic encephalopathy; DLVP, discontinuous low-voltage pattern; DHVP, discontinuous high-voltage pattern; SWC, sleep-wake cycling; TTNT, time to normal trace.

Imaging findings

Cranial ultrasound findings in the extremely preterm group were as follows: nine (39%) had normal cranial ultrasound, one (4.3%) had PIVH grade I, four (17.3%) had PIVH grade II, four (17.3%) had PIVH grade III, and three (13%) had PIVH grade IV. Three (13%) infants developed PVL and cranial ultrasound was not performed in two due to early premature death. MRI findings in infants with HIE are shown in Table 3. MRI was not performed in one of the patients due to premature death.

Interobserver agreements between radiologists regarding injury of the BGT, white matter, and cortical gray matter were: 75% ($\kappa = 0.636$; $p < 0.001$); 56.3% ($\kappa = 0.321$; $p = 0.029$), and 81.3% ($\kappa = 0.586$; $p < 0.001$), respectively. The observed agreement between radiologists regarding the PLIC was 100% ($\kappa = 1.0$; $p < 0.001$). Analyses of the normal/mild and moderate/severe injury groups of the BGT, white matter, and cortical gray matter were: 100% ($\kappa = 1.0$; $p < 0.001$); 87.5% ($\kappa = 0.589$; $p = 0.009$), and 93.8% ($\kappa = 0.764$; $p < 0.001$), respectively.

Discussion

This study was performed to evaluate the correlation between electroencephalic disorders and early outcome in a prospective cohort of infants at high risk for brain injury in a sample of the Brazilian population. The absence of SWC in the first 72 h of life and the presence of pathological trace or DLVP activity were associated with severe PIVH or

Table 3 MRI findings in infants with HIE.

	Total n (%)	Poor early outcome n (%)	Good early outcome n (%)
<i>Lesion in PLIC</i>			
Present	14(87.5%)	2(12.5%)	12(75%)
Absent	2(12.5%)	2(12.5%)	0(0%)
<i>Lesion in BGT</i>			
Absent	6(37.5%)	0(0%)	6(37.5%)
Mild	6(37.5%)	1(6.25%)	5(31.25%)
Moderate	2(12.5%)	2(12.5%)	0(0%)
Severe	2(12.5%)	2(12.5%)	0(0%)
<i>Lesion in white matter</i>			
Absent	6(37.5%)	0(0%)	6(37.5%)
Mild	7(43.75%)	2(12.5%)	5(31.25%)
Moderate	1(6.25%)	1(6.25%)	0(0%)
Severe	2(12.5%)	2(12.5%)	0(0%)
<i>Lesion in cortical gray matter</i>			
Absent	2(12.5%)	0(0%)	2(12.5%)
Mild	11(68.75%)	2(12.5%)	9(56.25%)
Moderate	1(6.25%)	1(6.25%)	0(0%)
Severe	2(12.5%)	2(12.5%)	0(0%)

MRI, magnetic resonance imaging; HIE, hypoxic-ischemic encephalopathy.

death in the neonatal period. Infants with poor early outcome more commonly changed from normal to pathological aEEG pattern than the opposite. These findings support the findings of Soubasi et al.,¹⁰ which showed a sensitivity of 89% and 70%, and specificity of 80% and 80.7% for background activity (pathological trace or DLVP) and absence of SWC, respectively, for the occurrence of adverse outcomes (PIVH grade III/IV, PVL, or death). A recent prospective cohort study demonstrated that preterm infants with severe cerebral lesions already manifest a maturational delay in the aEEG cyclic activity soon after birth.²¹ Others studies have also reported a significant positive correlation between the degree of abnormality of the aEEG and both USG abnormalities and poor neurodevelopmental prognosis.^{22,23}

In newborns with HIE, the presence of seizures (repetitive seizures or status epilepticus) patterns on aEEG/EEG traces and longer TTNT were associated with moderate to severe lesions observed on MRI and death. Earlier studies also found an association between electrographic seizures and severity of findings on cranial MRI.^{24,25} A retrospective review of continuous aEEG data from encephalopathic newborns treated with whole-body hypothermia found association between delay of aEEG background recovery as increasingly predictive of adverse early outcome over time in newborns being treated with therapeutic hypothermia.²⁶ Thoresen et al. have described that the recovery time to normal background pattern was the best predictor of poor outcome by the age at 2 years in neonates with birth asphyxia.²⁷ A recent meta-analysis showed that aEEG can be used as predictive and prognostic tool in preterm infants, with a specificity of 93% and sensitivity of 84%.²⁸

The present study showed a good interobserver agreement in MRI evaluations; however, should this be interpreted with caution due to the small sample size.

Strengths of this study included an independent and blinded radiographic review and the use of two-channel montage associated with the interpretation of raw EEG technique (aEEG/EEG), as it is known to increase the sensitivity and specificity of the method.^{29,30} Limitations of the study were the relatively small sample size, the lack of stratification by the use of antiepileptic drugs, sedation or other medications which might potentially influence the aEEG trace, and lack of knowledge regarding timing of brain injury, especially in the preterm group, in which injury could occur after the period of aEEG monitoring.

In conclusion, this study confirms that previous results are applicable to the studied population, and thus aEEG is a justified tool in the neonatal intensive care unit to monitor this group of neonates at high risk for brain injury.

Conflicts of interest

The authors declare no conflicts of interest.

References

1. Blencowe H, Lee AC, Cousens S, Bahalim A, Narwal R, Zhong N, et al. Preterm birth-associated neurodevelopmental impairment estimates at regional and global levels for 2010. *Pediatr Res.* 2013;74:17–34.
2. Lee AC, Kozuki N, Blencowe H, Vos T, Bahalim A, Darmstadt GL, et al. Intrapartum-related neonatal encephalopathy incidence and impairment at regional and global levels for 2010 with trends from 1990. *Pediatr Res.* 2013;74:50–72.
3. Leijser LM, de Bruïne FT, Steggerda SJ, van der Grond J, Walther FJ, van Wezel-Meijler G. Brain imaging findings in very preterm infants throughout the neonatal period: part I. Incidences and evolution of lesions, comparison between ultrasound and MRI. *Early Hum Dev.* 2009;85:101–9.
4. Shankaran S, Slovis TL, Bedard MP, Poland RL. Sonographic classification of intracranial hemorrhage. A prognostic indicator of mortality, morbidity, and short-term neurologic outcome. *J Pediatr.* 1982;100:469–75.
5. Watanabe K, Hayakawa F, Okumura A. Neonatal EEG: a powerful tool in the assessment of brain damage in preterm infants. *Brain Dev.* 1999;21:361–72.
6. Le Bihannic A, Beauvais K, Busnel A, de Barace C, Furby A. Prognostic value of EEG in very premature newborns. *Arch Dis Child Fetal Neonatal Ed.* 2012;97:F106–9.
7. Okumura A, Hayakawa F, Kato T, Kuno K, Watanabe K. Developmental outcome and types of chronic-stage EEG abnormalities in preterm infants. *Dev Med Child Neurol.* 2002;44:729–34.
8. Bowen JR, Paradisis M, Shah D. Decreased aEEG continuity and baseline variability in the first 48 hours of life associated with poor short-term outcome in neonates born before 29 weeks gestation. *Pediatr Res.* 2010;67:538–44.
9. Chalak LF, Sikes NC, Mason MJ, Kaiser JR. Low-voltage aEEG as predictor of intracranial hemorrhage in preterm infants. *Pediatr Neurol.* 2011;44:364–9.
10. Soubasi V, Mitsakis K, Sarafidis K, Griva M, Nakas CT, Drossou V. Early abnormal amplitude-integrated electroencephalography (aEEG) is associated with adverse short-term outcome in premature infants. *Eur J Paediatr Neurol.* 2012;16:625–30.
11. Toso PA, González AJ, Pérez ME, Kattan J, Fabres JG, Tapia JL, et al. Clinical utility of early amplitude integrated EEG in monitoring term newborns at risk of neurological injury. *J Pediatr (Rio J).* 2014;90:143–8.
12. al Naqeeb N, Edwards AD, Cowan FM, Azzopardi D. Assessment of neonatal encephalopathy by amplitude-integrated electroencephalography. *Pediatrics.* 1999;103:1263–71.
13. Toet MC, Hellström-Westas L, Groenendaal F, Eken P, de Vries LS. Amplitude integrated EEG 3 and 6 hours after birth in full term neonates with hypoxic-ischaemic encephalopathy. *Arch Dis Child Fetal Neonatal Ed.* 1999;81:F19–23.
14. Klebermass K, Olischar M, Waldhoer T, Fuiiko R, Pollak A, Weninger M. Amplitude-integrated EEG pattern predicts further outcome in preterm infants. *Pediatr Res.* 2011;70:102–8.
15. Wikström S, Pupp IH, Rosén I, Norman E, Fellman V, Ley D, et al. Early single-channel aEEG/EEG predicts outcome in very preterm infants. *Acta Paediatr.* 2012;101:719–26.
16. Sarnat HB, Sarnat MS. Neonatal encephalopathy following fetal distress. A clinical and electroencephalographic study. *Arch Neurol.* 1976;33:696–705.
17. Hellström-Westas L, Rosén I. Continuous brain-function monitoring: state of the art in clinical practice. *Semin Fetal Neonatal Med.* 2006;11:503–11.
18. Olischar M, Klebermass K, Kuhle S, Hulek M, Kohlhauser C, Rücklinger E, et al. Reference values for amplitude-integrated electroencephalographic activity in preterm infants younger than 30 weeks' gestational age. *Pediatrics.* 2004;113:e61–6.
19. Mizrahi EM. Acute and chronic effects of seizures in the developing brain: lessons from clinical experience. *Epilepsia.* 1999;40:S42–50, discussion S64–6.
20. Rutherford M, Ramenghi LA, Edwards AD, Brocklehurst P, Halliday H, Levene M, et al. Assessment of brain tissue injury after moderate hypothermia in neonates with hypoxic-ischaemic encephalopathy: a nested substudy of a randomised controlled trial. *Lancet Neurol.* 2010;9:39–45.

21. Natalucci G, Rousson V, Bucher HU, Bernet V, Hagmann C, Latal B. Delayed cyclic activity development on early amplitude-integrated EEG in the preterm infant with brain lesions. *Neonatology*. 2013;103:134–40.
22. Song J, Zhu C, Xu F, Guo J, Zhang Y. Predictive value of early amplitude-integrated electroencephalography for later diagnosed cerebral white matter damage in preterm infants. *Neuropediatrics*. 2014;45:314–20.
23. Benavente-Fernández I, Lubián-López SP, Jiménez-Gómez G, Lechuga-Sancho AM, Garcia-Alloza M. Low-voltage pattern and absence of sleep-wake cycles are associated with severe hemorrhage and death in very preterm infants. *Eur J Pediatr*. 2015;174:85–90.
24. Shah DK, Wusthoff CJ, Clarke P, Wyatt JS, Ramaiah SM, Dias RJ, et al. Electrographic seizures are associated with brain injury in newborns undergoing therapeutic hypothermia. *Arch Dis Child Fetal Neonatal Ed*. 2014;99:F219–24.
25. Glass HC, Nash KB, Bonifacio SL, Barkovich AJ, Ferriero DM, Sullivan JE, et al. Seizures and magnetic resonance imaging-detected brain injury in newborns cooled for hypoxic-ischemic encephalopathy. *J Pediatr*. 2011;159:731–5.e1.
26. Massaro AN, Tsuchida T, Kadom N, El-Dib M, Glass P, Baumgart S, et al. aEEG evolution during therapeutic hypothermia and prediction of NICU outcome in encephalopathic neonates. *Neonatology*. 2012;102:197–202.
27. Thoresen M, Hellström-Westas L, Liu X, de Vries LS. Effect of hypothermia on amplitude-integrated electroencephalogram in infants with asphyxia. *Pediatrics*. 2010;126:e131–9.
28. Lin B, Zhang P, Cheng G, Zhou W, Wang L. Meta-analysis of prognostic tests in neonates over 35-week gestational age with hypoxic-ischemic encephalopathy. *Zhonghua Yi Xue Za Zhi*. 2014;94:115–21.
29. Lavery S, Shah DK, Hunt RW, Filan PM, Doyle LW, Inder TE. Single versus bihemispheric amplitude-integrated electroencephalography in relation to cerebral injury and outcome in the term encephalopathic infant. *J Paediatr Child Health*. 2008;44:285–90.
30. Mastrangelo M, Fiocchi I, Fontana P, Gorgone G, Lista G, Belcastro V. Acute neonatal encephalopathy and seizures recurrence: a combined aEEG/EEG study. *Seizure*. 2013;22:703–7.